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TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371

XI/P5934USO

U.S. APPLICATION NO. (if known, see 37 CFR 1.5)

INTERNATIONAL APPLICATION NO.

PCT/JP96/03344

INTERNATIONAL FILING DATE

14 November 1996

PRIORITY DATE CLAIMED

14 November 1995

TITLE OF INVENTION

MATERIAL FOR NEGATIVE ELECTRODE OF LITHIUM SECONDARY BATTERY, ...

APPLICANT(S) FOR DO/EO/US

MORITA, Koichi et al.

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☐ have not been made and will not be made.
8. ☒ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☒ A translation of the annexes to the International Preliminary Examination Report under PCT Article 34 (35 U.S.C. 371(c)(5)). - (2) SETS - 11/17/97 + 11/12/98.

Items 11. to 16. below concern document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
 12. ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
 13. ☒ A **FIRST** preliminary amendment.
☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
 14. ☐ A substitute specification.
 15. ☐ A change of power of attorney and/or address letter.
 16. ☒ Other items or information:
 - a. Customer Number/Change of Address Letter
 - b.
 - c.
 - d.
- 16e. ☐ A copy of the Notification of Missing Requirements under 35 USC 371.
- 16f. ☐ In the event that a petition for extension of time is required to be submitted herewith, and in the event that a separate petition does not accompany this response, applicant hereby petitions under 37 CFR 1.136(a) for an extension of time as many as are required to render this submission timely. Any fee is authorized below in 17(c).

Date: 14 May 1998

17. ☒ The following fees are submitted:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5))

Search Report has been prepared by the EPO (or JPO) \$ 930.00

International preliminary examination fee paid to USPTO (37 CFR 1.482) \$ 720.00

No international preliminary examination fee paid to USPTO (37 CFR 1.482)
but international search fee paid to USPTO (37 CFR 1.445(a)(2)) \$ 790.00Neither international preliminary examination fee (37 CFR 1.482) nor
international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$1070.00International preliminary examination fee paid to USPTO (37 CFR 1.482)
and all claims satisfied provisions of PCT Article 33(2)-(4) \$ 98.00

ENTER APPROPRIATE BASIC FEE AMOUNT =

CALCULATIONS PTO USE ONLY

\$ 930

Surcharge of \$130.00 for furnishing the oath or declaration later than ☐ 20 ☐ 30
months from the earliest claimed priority date (37 CFR 1.492(e)).

\$

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total claims	37 - 20 =	17	X \$22.00
Independent claims	05 - 3 =	02	X \$82.00
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$270.00

\$ 374

\$ 164

\$

TOTAL OF ABOVE CALCULATIONS =

\$ 1468

Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity Statement
must also be filed (Note 37 CFR 1.9, 1.27, 1.28).

\$

SUBTOTAL =

\$ 1468

Processing fee of \$130.00 for furnishing the English translation later than ☐ 20 ☐ 30
months from the earliest claimed priority date (37 CFR 1.492(f)).

\$

+

TOTAL NATIONAL FEE =

\$ 1468

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be
accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property

\$

+

TOTAL FEES ENCLOSED =

\$ 1508

Amount to be:
refunded

\$

charged

\$

a. ☒ A check in the amount of \$ 1508 to cover the above fees is enclosed.b. ☐ Please charge my Deposit Account No. 12-0555 in the amount of _____ to cover the above fees.c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit
overpayment to Deposit Acct. No. 12-0555.Note: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR
1.137(a) or (b) must be filed and granted to restore the application to pending status.Please address all correspondence to Thomas P. Sarro at the address (below) of
Customer Number: 00881.

SEND ALL CORRESPONDENCE TO:

LARSON & TAYLOR

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B. Aaron Schulman

Name

31877

Registration

Date: 14 May 1998

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Patent

In re application of **MORITA, et al****88 Rec'd PCT/PTO 14 MAY 1998**

New Application

) Atty's Dckt: XI/P5934US0

Filed: On even date herewith

) Application Branch

For: MATERIAL FOR NEGATIVE ELECTRODE OF LITHIUM SECONDARY...

PRELIMINARY AMENDMENTHon. Assistant Commissioner of Patents
Washington, D.C. 20231

S I R:

Preliminary to the examination thereof, please amend the above-identified application as follows:

IN THE CLAIMS:

Claim 3, lines 1 and 2, delete "or 2".

Claim 4, lines 1 and 2, delete "any of claims 1-3", and insert therefor --claim 1--.

Claim 5, lines 1 and 2, delete "any of claims 1-4", and insert therefor --claim 1--.

Claim 6, lines 1 and 2, delete "any of claims 1-5", and insert therefor --claim 1--.

Claim 9, line 2, delete "or 8".

Claim 10, line 2, delete "any of claims 7-9", and insert therefor --claim 7--.

Claim 11, line 2, delete "any of claims 7-10", and insert therefor --claim 7--.

Claim 12, line 2, delete "any of claims 7-11", and insert therefor --claim 7--.

Claim 15, line 2, delete "or 14".

Claim 16, line 2, delete "any of claims 13-15", and insert therefor --claim 13--.

Claim 17, line 2, delete "any of claims 13-16", and insert therefor --claim 13--.

Claim 18, line 2, delete "any of claims 13-17", and insert therefor --claim 13--.

Claim 19, line 2, delete "any of claims 13-18", and insert therefor --claim 13--.

Claim 20, line 2, delete "any of claims 13-19", and insert therefor --claim 13--.

Claim 21, line 2, delete "any of claims 13-20", and insert therefor --claim 13--.

Claim 22, line 2, delete "any of claims 13-21", and insert therefor --claim 13--.

Claim 23, line 3, delete "any of claims 1-6", and insert therefor --claim 1--.

Claim 24, line 3, delete "any of claims 1-6", and insert therefor --claim 1--.

Claim 25, line 3, delete "any of claims 1-6", and insert therefor --claim 1--.

Claim 26, line 3, delete "any of claims 1-6", and insert therefor --claim 1--.

09/068592-054199

Claim 27, line 2, delete "any of claims 23-26", and insert therefor --claim 23--.

Claim 28, line 2, delete "any of claims 1-12", and insert therefor --claim 1--.

Claim 29, line 3, delete "any of claims 1-12", and insert therefor --claim 1--.

Claim 30, line 2, delete "or 29".

Claim 37, line 2, delete "any of claims 32-36", and insert therefor --claim 32--.

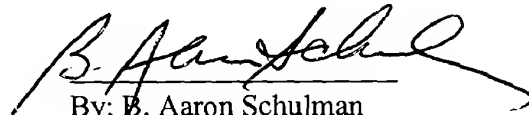
REMARKS

The above amendments are being made in order to place the application in better condition for examination and to reduce the filing fee.

Favorable consideration is respectfully requested.

Respectfully submitted,

Date: May 14, 1998


By: B. Aaron Schulman
Registration No. 31877

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PCT

特許協力条約に基づいて公開された国際出願

<p>(51) 国際特許分類 C01B 31/02, 31/04, H01M 4/58, 4/02, 10/40, 4/02</p>	<p>A1</p>	<p>(11) 国際公開番号 WO97/18160</p> <p>(43) 国際公開日 1997年5月22日 (22.05.97)</p>
<p>(21) 国際出願番号 PCT/JP96/03344</p> <p>(22) 国際出願日 1996年11月14日 (14.11.96)</p> <p>(30) 優先権データ 特願平7/295462 1995年11月14日 (14.11.95)</p> <p>(71) 出願人 (米国を除くすべての指定国について) 大阪瓦斯株式会社(OSAKA GAS COMPANY LIMITED)[JP/JP] 〒541 大阪府大阪市中央区平野町四丁目1番2号 Osaka, (JP)</p> <p>(71) 出願人 (米国と日本を除くすべての指定国について) シャープ株式会社(SHARP CORPORATION)[JP/JP] 〒545 大阪府大阪市阿倍野区長池町22番22号 Osaka, (JP)</p> <p>(72) 発明者：および</p> <p>(75) 発明者／出願人 (米国についてのみ) 森田浩一(MORITA, Koichi)[JP/JP] 〒573 大阪府枚方市走谷1-25-47-203 Osaka, (JP) 藤原裕己(FUJIWARA, Hiromi)[JP/JP] 〒594 大阪府和泉市いぶき野1丁目18番2号 Osaka, (JP) 中川喜照(NAKAGAWA, Yoshiteru)[JP/JP] 〒639-11 奈良県大和郡山口市小泉町370 Nara, (JP) 勝浦将光(KATSUURA, Masamitsu)[JP/JP] 〒554 大阪府大阪市此花区西島6-3-4-506 Osaka, (JP) 松好弘明(MATSUYOSHI, Hiroaki)[JP/JP] 〒577 大阪府東大阪市金岡2-14-2 Osaka, (JP)</p>	<p>西村直人(NISHIMURA, Naoto)[JP/JP] 1 〒639-21 奈良県北葛城郡新庄町萱192-1 Nara, (JP) 佃 至弘(TSUKUDA, Yoshihiro)[JP/JP] 〒545 大阪府大阪市阿倍野区王子町4-1-9-306 Osaka, (JP) 湊 和明(MINATO, Kazuaki)[JP/JP] 〒536 大阪府大阪市城東区鳴野西5-1-2-1308 Osaka, (JP) 見立武仁(MITATE, Takchito)[JP/JP] 〒635 奈良県大和高田市野口65-4 Nara, (JP) 山田和夫(YAMADA, Kazuo)[JP/JP] 〒639-21 奈良県北葛城郡新庄町定田500-4 Nara, (JP) 米田哲也(YONEDA, Tetsuya)[JP/JP] 〒518-04 三重県名張市つづじが丘北9番町17番地 Mie, (JP)</p> <p>(74) 代理人 弁理士 三枝英二, 外(SAEGUSA, Eiji et al.) 〒541 大阪府大阪市中央区道修町1-7-1 北浜TNKビル Osaka, (JP)</p> <p>(81) 指定国 CA, CN, JP, KR, US, 欧州特許 (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).</p> <p>添付公開書類 国際調査報告書 補正書・説明書</p>	
<p>(54) Title: CATHODE MATERIAL FOR LITHIUM SECONDARY BATTERY, PROCESS FOR MANUFACTURING THE SAME, AND SECONDARY BATTERY USING THE SAME</p>		
<p>(54) 発明の名称 リチウム二次電池用負極材料およびその製造方法、ならびにそれを用いた二次電池</p>		
<p>(57) Abstract</p> <p>A carbon material for a lithium secondary battery, in which the edge parts of the crystals of a carbon material which is the core material of the cathode material are partially or entirely coated with a carbon material and the crystals have nearly spherical or elliptic shapes. There are also provided a process for manufacturing the carbon material, and a lithium secondary battery using the carbon material as a component.</p>		

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SPECIFICATION

MATERIAL FOR NEGATIVE ELECTRODE OF LITHIUM SECONDARY
BATTERY, METHOD FOR PRODUCTION THEREOF AND LITHIUM
SECONDARY BATTERY USING THE SAME

5

TECHNICAL FIELD

The present invention relates to a carbon
material and a method for production thereof, in
particular, a carbon powder whose surface is coated with
ingredients of heavy oil and a method for production
10 thereof. Specifically, the invention relates to a carbon
material, useful as material for negative electrode of
lithium secondary battery, a method for production
thereof and lithium secondary battery using such a carbon
material.

15

BACKGROUND ART

Recently, miniaturization and weight-saving of
electronic equipments, information related equipments and
like portable-type equipments (hereinafter "portable
equipment") have been progressed significantly, which
20 makes secondary batteries driving said equipments very
important parts. Lithium secondary batteries have
lightness in weight and high energy density so that
lithium secondary batteries are regarded as promising
driving power source of portable equipment. Research and
25 development thereof are actively progressed. When

lithium metal is used as negative electrode, dendrite formed and grown on metal lithium by repeating a charge and discharge cycle causes an internal short circuit, which makes production of lithium secondary battery
5 difficult. Although use of lithium alloys, such as a lithium aluminium alloy in place of lithium metal is proposed, batteries using the alloy causes segregation of alloy in the course of charge and discharge cycle or deep charge and discharge. Consequently, batteries
10 maintaining sufficient characteristics for a long period of time can not be obtained.

Batteries using carbon materials as host material of negative electrode to utilize an intercalation and deintercalation reaction of lithium ion
15 are proposed, developed and come in practice. Lithium secondary batteries applying carbon materials to a negative electrode is superior in cycle characteristics, safety, etc.

Carbon materials have a variety of structures
20 and forms from graphite to amorphous carbon. Performance of electrode is greatly influenced by physical properties and microstructure thereof formed by hexagonal net face of carbon. A variety of carbon materials whose physical properties and microstructure are specified are,
25 therefore, proposed. Presently used materials for

negative electrode of lithium secondary battery are roughly classified into carbon-series materials calcined at about 1,000 °C and graphite-series materials calcined at about 2,800 °C. The former has an advantages of low reactivity with electrolyte leading to difficulty of electrolyte decomposition and a drawback of great change of electric potential with release of lithium ion. In contrast, the latter has an advantage of small change of electric potential with release of lithium ion and a drawback of reaction with electrolyte leading to decomposition of electrolyte and destruction of carbon materials (J. Electrochem. Soc. 117, 222 (1970)). As a result, the latter causes problems of decreased efficiency of charge and discharge, decreased cycle characteristics and decreased safety of battery. It is reported that graphite-series materials may be used, when specific electrolytes are used (J. Electrochem. Soc. 137, 2009(1990)). For the purpose of producing battery, however, limitation of electrolyte has a drawback of restraint of improvement of temperature characteristics and cycle characteristics of battery by the type of electrolyte.

To solve the problems, JP-A-4-368778, JP-A-4-370662, JP-A-5-94838 and JP-A-5-121066 proposed carbon materials produced by coating a surface of graphite

particles with low-crystalline carbon. These surface-modified carbon materials are effective to increase battery capacity and to improve cycle characteristics through inhibition of electrolyte decomposition.

5 According to techniques described in JP-A-4-368778, carbon coating layer formed in gas phase do not cause fusing and aggregation of carbon particles leading to generation of materials with high performance. However, the materials have practical problems in cost
10 and mass production.

 JP-A-4-370662, JP-A-5-94838 and JP-A-5-121066 disclose a method of coating carbon in liquid phase which is advantageous in cost and mass production. However, a simple combination of mixing graphite particles and
15 organic compounds in liquid phase with calcination thereof generates fused and aggregated carbon particles which are necessary to be powdered, causing drawbacks of exposure of generated active surface of graphite by
20 powdering fused and aggregated carbon particles, contamination during powdering and complexity of production steps.

DISCLOSURE OF THE INVENTION

 It is a primary object of the invention to obtain a lithium secondary battery which is superior in
25 properties such as cycle characteristics and safety by

manufacturing a negative electrode using carbon materials which are free of selectivity and/or restraint with respect to electrolytes, and small in change of electric potential with release of lithium ion.

5 The inventors conducted extensive research to solve or decrease said prior art problems and found that carbon materials whose core material is uniformly covered with pitch can be produced by dipping particle-like carbon materials to be core material (hereinafter
10 referred to as "core carbon material" or "carbon material to be core material" or simply "core material") into raw material for coat-forming carbon material (for example, tar, pitch and like coal heavy oil or petroleum heavy oil; hereinafter may be referred to as simply "heavy oil
15 etc."), followed by taking specific means to separate core material from heavy oil etc. It is found that the two-layered carbon material particles thus obtained have a spheric or ellipsoidal or similar shape, or a shape that edge parts of carbon crystal are rounded. As a
20 result of measurement by a BET method, it is found that specific surface area of the particles are smaller than core carbon materials before treatment showing that pores concerning specific surface area determined by a BET method are filled in a certain way.

25 According to the invention, coated carbon

material particles characterized in that carbon materials derived from heavy oil etc. are partially or entirely adhered to edge and basal plane of carbon materials to be core material, or, part or whole of the edge and basal plane are coated by the carbon material to be substantially spheric or ellipsoidal shape are provided. With respect to the carbon materials, pores concerning specific surface area determined by a BET method are filled by adhesion or coating of carbon derived from heavy oil etc. thereto, and the carbon materials have a specific surface area of $5\text{m}^2/\text{g}$ or less (preferably about $1\text{-}5\text{m}^2/\text{g}$).

According to the invention, carbon materials to be core material are highly crystalline graphite material having a mean interplanar spacing (d_{002}) of (002) plane of 0.335-0.340 nm, a thickness of crystallite size in direction of (002) plane (L_c) of at least 10 nm (preferably at least 40 nm), a thickness of crystallite size in direction of (110) plane (L_a) of at least 10 nm (preferably at least 50 nm) determined by X-ray wide-angle diffractometry.

The carbon material of the invention is characterised by lower crystallinity of carbon materials adhered to or coated on the surface of core material (hereinafter may be referred to as "carbon material for

coating formation") than crystallinity of said core material.

The carbon material of the invention has a true specific gravity of 1.50 to 2.26g/cm³.

5 A lithium secondary battery with high capacity and safety may be obtained by using such carbon material as material of negative electrode of lithium secondary battery.

10 Said coated carbon material of the invention may be produced as follows. The carbon material to be core material is dipped into tar, pitch and like coal heavy oil or petroleum heavy oil, preferably at about 10-300 °C for coating with heavy oil etc., followed by separating the coated core carbon material from heavy
15 oil, adding an organic solvent to the coated carbon material thus separated for washing preferably at about 10-300 °C, and then drying the material.

20 Furthermore, the invention provides a method for producing a carbon material comprising calcining for carbonization the carbon material coated with heavy oil etc. obtained according to the above-mentioned procedure and a method for producing a carbon material comprising calcining for graphitization the carbon material coated
25 by heavy oil etc. obtained according to the above-mentioned procedure.

With respect to the invention, the carbon material obtained according to said manufacturing method comprises preferably 10% or less of particles having a diameter of 1 μm or less as volume-based integrated value, wherein said diameter is determined by laser diffraction-type particle size distribution.

According to the invention, tar or pitch modified by removing at least part of primary QI to reduce a remaining primary QI of 3% or less (preferably 1% or less) is preferable.

Furthermore, the invention provides a material for negative electrode of lithium secondary battery characterized in that the material comprises said carbonized or graphitized carbon materials; a negative electrode for lithium secondary battery using the material for negative electrode; and also a non-aqueous lithium secondary battery and a solid electrolyte secondary battery using the negative electrode.

With respect to the invention, "nearly spheric or ellipsoidal" carbon materials include carbon materials free of sharp edge by adhesion of carbon ingredients from heavy oil etc. to edge and basal plane of carbon material to be core material partially or entirely, and shapes of carbon material particles to be core material observed by SEM are maintained. The carbon material may be

efficiently produced by the method of the invention free of grinding step. However, the material is not limited to the material produced by the method of the invention.

With respect to the invention, carbon materials
5 whose "pores concerning specific surface area determined by a BET method are coated and filled by raw material for coat-forming carbon material, i.e., carbon materials from tar, pitch and like coal heavy oil or petroleum heavy oil adhered thereon" include carbon materials whose pores
10 concerning specific surface area determined by a BET method is filled at least part thereof by calcined product of raw material for coat-forming carbon material (hereinafter referred to as coat-forming carbon material). Specifically, it is not necessary for the
15 pores to be completely filled by carbon materials derived from heavy oil etc. For example, carbon materials whose adjacent entry ports are filled are included. Such conditions of carbon materials may be confirmed by reduction of specific surface area determined by a BET
20 method .

The carbon materials obtained according to the invention include 4 combinations, i.e., low-crystallinity carbon material + low-crystallinity carbon material; low-crystallinity carbon material + high-crystallinity carbon
25 material; high-crystallinity carbon material + low-

crystallinity carbon material; and high-crystallinity carbon material + high-crystallinity carbon material. In all cases, an effect on decrease of electrolyte degradation is exerted.

5 With respect to the invention, low-crystallinity carbon materials means "carbon which may not be graphite crystals by a treatment necessary for graphitization (eg. treatment at elevated temperature)". Such carbon is usually referred to as hard carbon. On
10 the other hand, high-crystallinity carbon material means "carbon which becomes graphite crystals by a treatment for graphitization". Such carbon is usually referred to as soft carbon.

 According to the invention, the following 8
15 carbon materials are obtained depending on a combination of core material and outer carbon material (which may be referred to as "coat-forming carbon material", "carbon material for surface-modification", "covering material") and on final calcination temperature. Examples are:

- 20 (1) carbonized carbon materials comprising core material consisting of low-crystallinity carbon materials and coat-forming carbon materials consisting of low-crystallinity carbon materials;
- (2) carbonized carbon materials comprising core material
25 consisting of low-crystallinity carbon materials and

coat-forming carbon materials consisting of high-crystallinity carbon materials;

(3) graphitized carbon materials comprising core material consisting of low-crystallinity carbon materials and

5 coat-forming carbon materials consisting of low-crystallinity carbon materials;

(4) graphitized carbon materials comprising core material consisting of low-crystallinity carbon materials and coat-forming carbon materials consisting of high-

10 crystallinity carbon materials;

(5) carbonized carbon materials comprising core material consisting of high-crystallinity carbon materials and coat-forming carbon materials consisting of low-crystallinity carbon materials;

15 (6) carbonized carbon materials comprising core material consisting of high-crystallinity carbon materials and coat-forming carbon material consisting of high-crystallinity carbon materials;

(7) graphitized carbon materials comprising core material
20 consisting of high-crystallinity carbon materials and coat-forming carbon material consisting of low-crystallinity carbon materials;

(8) graphitized carbon materials comprising core material consisting of high-crystallinity carbon materials and

25 coat-forming carbon material consisting of high-

crystallinity carbon materials.

According to the invention, carbon materials for secondary batteries having small specific surface area and good charge and discharge properties may be efficiently produced by coating core material with outer carbon materials. In particular, carbon materials for battery with excellent charge and discharge properties may be obtained by a combination of core material and covering material shown in (5), (6) and (7), and also carbon materials for battery with small specific surface area and improved safety may be obtained by a combination of core material and covering material shown in (1), (2), (3), (4) and (8).

According to the invention, as carbon material to be core material, one or more of particle-like (scaly or massive, fibrous, whisker-like, spheric, shattered etc.) natural graphite, artificial graphite, mesocarbon microbeads, mesophase pitch powder, isotropic pitch powder, resin and carbonized and graphitized products thereof. In particular, scaly and massive natural graphite and artificial graphite which are very inexpensive are preferable from the viewpoint of cost. The carbonized and graphitized products of mesocarbon microbeads (MCMB) having very small specific surface area leading to obtaining material having smaller specific

surface area are preferable from the viewpoint of safety of secondary battery, if the products are used as core material.

Carbon materials to be core material are more preferably, 0.335-0.340 nm in mean interplanar spacing (d002) of (002) plane, at least 10nm (preferably at least 40 nm) in thickness of crystallite size in direction of (002) plane (Lc), at least 10 nm (preferably at least 50 nm) in thickness of crystallite size in direction of (110) plane (La) determined by X-ray wide-angle diffractometry, and 0.5 or less (preferably 0.4 or less) in ratio of peak strength around 1360cm^{-1} to peak strength around 1580cm^{-1} (hereinafter referred to as R value) determined by Raman spectroscopy with argon laser.

When the mean interplanar spacing is more than 0.340 nm, or Lc and La are smaller than 10nm, or R value is more than 0.5, crystallinity of carbon materials is insufficient, and covering carbon materials produced therefrom are not preferable because of insufficient capacity at low electric potential near dissolution and deposition of lithium (0-300mV on the potential vs. Li/Li^+).

Particle size distribution of carbon materials to be core material is preferably about 0.1-150 μm .

Since particle size of final product containing coat-

forming carbon material derived from heavy oil etc.
substantially depends on particle size of carbon material
to be core material, particle size of the final product
is substantially specified by particle size of core
5 material. When particle size of core material is less
than 0.1 μm , internal short circuit is likely to be
caused through pores of separator of battery, thereby not
preferable. On the other hand, when particle size of
core material is more than 150 μm , uniformity of
10 electrode, packing density of active material and
handling properties during steps of production of
electrode are decreased, thereby not preferable.

A weight ratio of coat-forming carbon material
derived from heavy oil etc., that is, coat-forming carbon
15 material/(core carbon material + coat-forming carbon
material) (hereinafter the ratio is referred to as
"covering ratio") is preferably more than 0 and up to
0.3, more preferably 0.01-0.2. In this case, thickness
of coat-forming carbon is about 0.01-10 μm , more
20 preferable thickness is about 0.05-5 μm .

When a covering ratio exceeds 0.3, to ensure
sufficient capacity of battery produced therefrom becomes
difficult. The amount of coat-forming carbon is
determined as quinoline soluble matter by solvent
25 analysis of carbon components from heavy oil etc. which

cover surface of core material before calcination.

Thickness of coat-forming carbon materials is determined by measuring a central particle size (D50) of carbon material to be core material before coating and a central
5 particle size (D50) of pitch-coated carbon materials before calcination with a laser diffraction particle size analyzer, followed by calculating the thickness using the equation : $\{(\text{particle size after coating}) - (\text{particle size of raw material before coating})\} / 2$ based on the
10 assumption that carbon materials are spheric and that shape of coat layer comprising pitch components is maintained after calcination.

According to the invention, a combination that a coat-forming carbon material on the surface has lower
15 crystallinity than a core carbon material is preferable. Furthermore, a mean interplanar spacing (d002) of (002) plane of 0.335-0.340 nm, a thickness of crystallite size in direction of (002) plane (Lc) of at least 10 nm (preferably at least 40 nm) and a thickness of
20 crystallite size in direction of (110) plane (La) of at least 10 nm (preferably at least 50 nm) determined by X-ray wide-angle diffractometry, and 0.5 or more (preferably about 0.5-1.5) determined by Raman spectroscopy with argon laser are preferable. The
25 interplanar spacing and R value are general index of

crystallinity of graphite. From the nature of the measuring methods, X-ray diffractometry reflects bulk properties on determined value of physical property, on the other hand, Raman spectrometry reflects physical properties of surface of material. Specifically, materials which meet said physical properties mean that the materials have high crystallinity as bulk property and low crystallinity as surface thereof. When R value of material after calcination is less than 0.5, selectivity of solvent thereof is not completely removed because of high surface crystallinity. When mean interplanar spacing (d002) is outside of the range of 0.335-0.340 nm, change of electric potential with intercalation and deintercalation of lithium ion become large, thereby not preferable.

A true density of the coated carbon materials with two-layer structure obtained is about 1.50-2.26g/cm³, preferably about 1.8-2.26g/cm³, more preferably about 2.0-2.26g/cm³. When electrode is produced using material with low true density, obtaining battery with high capacity is difficult, since increase of density of active substance in electrode is not possible even if the material has superior properties per unit weight.

Coated carbon materials preferable have a

particle size distribution ranging from 0.1 to 150 μm .
Within the range of particle size distribution, the
material preferably has 10% or less of particles having a
diameter of 1 μm or less based on volume thereof. When
5 content of particles with a diameter of 1 μm is more than
10% based on volume thereof, battery properties are
decreased due to increase of specific surface area,
thereby not preferable.

The coated carbon materials in powder form
10 obtained according to the invention may be subjected to
mold fill-out, pressure molding and calcination to obtain
a carbon block or a graphite block with homogenous
composition.

Examples of raw material of coat-forming carbon
15 material are naphthalene, phenanthrene, acenaphthylene,
anthracene, triphenylene, pyrene, chrysene, perylene and
like aromatic hydrocarbons, tar or pitch obtained by
polycondensation thereof under pressure with heat, or,
tar, pitch, asphalt and oils containing a mixture of said
20 aromatic hydrocarbons as main component which may be
derived from petroleum oil and coal oil. In the
specification, the raw material of coat-forming carbon
material may be simply referred to as "(petroleum or
coal) heavy oil, etc.". Furthermore, a variety of
25 thermoset resins may be used as coat-forming raw

material, although they are disadvantageous in cost.

When coal heavy oil is used, tar or pitch having 3% or less (preferably 1% or less) of primary QI produced by removing at least part of primary QI which exists in raw material is preferably used. The primary QI means free carbon essentially included in coal tar. The primary QI which exists in raw material inhibits carbonization by calcination and is contaminated in the final product as spheric carbon particles having a diameter of about 1 μ m, which may result in introduction of problems in manufacturing process of electrode or decrease of electrode properties.

In general, heavy oil is solid at ordinary room temperature and may be softened and melted by heating. The temperature at which material become softening is referred to as softening point (SP). In order to specify quality of heavy oil, insoluble matter in toluene determined by solvent fractionation with toluene is usually used. Those are typical indications to specify heavy oil. According to the invention, optional indication may be suitably selected to specify quality of heavy oil.

According to the invention, carbon materials to be core material is mixed with heavy oil etc. and stirred. Stirring methods are not specifically limited

to, but include mechanically stirring methods using ribbon mixer, screw-type kneader, universal mixer and the like.

Conditions for stirring treatment (temperature and time) are suitably selected in compliance with components of raw material (core material and coat-forming heavy oil) and viscosity of mixture. The conditions are usually about 10-300°C, more preferably about 50-200°C. Time period may be determined to make viscosity of mixture of 5000 Pa·s or less. The thickness of coating layer of coat-forming raw material (which hereinafter may be referred to as simply coating layer) may be controlled by adjusting temperature and time period during a stirring treatment. Higher temperature and/or shorter time lead to a thinner coating layer. In contrast, lower temperature lead to a thicker coating layer. Insufficient stirring causes ununiform coating layer, thereby not preferable. In general, stirring time does not adversely affect properties of product. However, too long stirring time practically decreases productivity, thereby not preferable. Time period may be suitably selected.

Atmosphere during stirring may be under any of atmospheric pressure, application of pressure and reduced pressure. Stirring under reduced pressure improves

conformability of core material with heavy oil, thereby preferable.

According to the invention, a plural of mixing and stirring processes may be conducted for the purpose of increased conformability of core material with heavy oil, uniform thickness of coating layer and thicker coating, if necessary. Before the following washing step, the coated core material may be separated and then subjected to washing step.

Subsequently, the carbon materials coated by heavy oil etc. thus obtained is subjected to a washing step. Examples of organic solvent used for washing are toluene, quinoline, acetone, hexane, benzene, xylene, methylnaphthalene, alcohols, solvents from coal oil and petroleum oil. Among them, toluene, quinoline, acetone, benzene, xylene, methanol, gas oil and middle oil from coal and petroleum are more preferable. Heavy oil components in coating layer may be controlled by imparting insoluble matters in washing solvent to coating layer, if organic solvents are suitably selected.

Washing temperature is determined according to finally obtainable coated carbon materials, in particular, properties of surface of coating layer. The temperature is not specifically limited, but is preferably about 10-300 °C.

A proportion of solid matter { = core material + coating layer or impregnation layer (hereinafter simply referred to as coating layer)) and organic solvent during washing is preferably 1 : 0.1-10 by weight ratio.

5 In the washing step, thickness and remaining heavy oil components of coating layer may be controlled by selecting the type of solvent, washing time and washing temperature. For example, coating layer become thin by a suitable combination of solvent with high
10 detergency and elevating washing temperature. On the other hand, thickness of coating layer become thick by a suitable combination of solvent with low detergency and decrease of washing temperature. Washing time are selected according to said conditions.

15 Subsequently, separation step of coated carbon materials from organic solvent may be conducted according to centrifugation, press filtration, gravity settling and like techniques. Temperature during separation is usually about 10-300°C.

20 Drying of separated coated carbon materials is usually conducted about 100-400°C.

 The dried coated carbon materials maintain pitch components on the surface of core material particles leading to no fusing and aggregation of
25 particles.

Said coated carbon materials thus dried are then calcined. For carbonization of coated carbon materials, calcination may be carried out at temperature of about 600-2,000°C, preferably 900-1,300°C. For
5 graphitization, calcination may be carried out at temperature of about 2,000-3,000°C, preferably about 2,500-3,000°C.

In order to maintain low crystallinity during calcination at elevated temperature in conditions of
10 carbonization or graphitization, coated heavy oil layer may be subjected to a graphitization retardation treatment using oxidative gases such as oxygen, ozone, carbon monoxide and sulfur oxide before calcination of coated carbon materials, followed by calcined at elevated
15 temperature. For example, highly crystallizable coating layer is formed on highly crystallizable core material, and then oxidation treatment is conducted to convert a coating layer into lower crystallizable carbon. In contrast, highly crystallizable coating layer may be
20 maintained without said oxidation treatment. Such oxidation treatment is conducted before calcination of coated carbon materials. The carbon materials thus obtained is useful as material for negative electrode of lithium secondary battery.

25 Examples of atmosphere during calcination of

coated carbon materials are reducing atmosphere, inert gas flow, inert gas in closed system, vacuum condition and like non-oxidative atmosphere.

Irrespective of calcination temperature, a rate of elevation of temperature is selected from about 1-300°C/hr. Calcination time is about 6 hours to 1 month. Elevation of temperature may be conducted step-by-step in compliance with thickness of coating layer.

Vacuum calcination is preferably carried out with maintaining reduced pressure at a temperature from normal temperature to the highest temperature, or, at a suitable temperature range (preferably more than 500°C). Vacuum calcination is effective to remove surface functional groups of coated carbon materials leading to reduction of non-reversible capacity of battery.

In general, a rapid rate of elevating temperature is expected to improve productivity, and a slow rate of elevating temperature (up to 10°C/hr) is expected to form a densified coating layer. Temperature profile during elevation of temperature and calcination may be in a variety of manners such as linear elevation of temperature and stepwise elevation of temperature by holding temperature for a constant period of time.

When the carbon materials thus obtained whose surface is coated by coat-forming carbon materials are

applied to negative electrode of lithium secondary battery, decomposition of electrolyte and destruction of carbon materials are inhibited due to low reactivity thereof with organic solvent of electrolyte solution. As
5 a result, the battery has advantages of improved charge and discharge efficiency and safety. In general, graphite material has an outside-oriented active edge plane of crystal lattice so that graphite material is likely to react with electrolyte. According to the
10 invention, since the active edge plane of crystal lattice is covered by pitch components whose basal plane, i.e., condensed polycyclic network of carbon, orientates outside, reaction with organic solvent of electrolyte will be inhibited.

15 According to the invention, since the amount of coating heavy oil on the surface of carbon materials and thickness of coating layer may be controlled by adjusting temperature and time of dipping carbon materials to be core material into heavy oil etc., or, the type of
20 organic solvent and washing conditions (time and temperature) for washing coated carbon materials, carbon materials whose surface is covered by pitch components may be produced, wherein said pitch components have basal plane, i.e., condensed polycyclic network of carbon,
25 which orientates in a direction of surface of carbon

materials.

Furthermore, with respect to the coating of core material surface, orientation of the basal plane in the direction of surface of carbon material is maintained during carbonization or following graphitization of these carbon materials. When the carbon material is applied to negative electrode of lithium secondary battery, since the carbon materials is unlikely to react with organic solvent, decomposition of electrolyte and destruction of carbon materials will not occur. Consequently, superior effects, such as high charge and discharge efficiency of battery and good safety of battery, are exerted.

In order to produce a lithium secondary battery of the invention, particle size of the coated carbon materials thus obtained is optionally adjusted by treatments such as dispersion and classification to be material for electrode.

Electrode is produced by mixing the carbon materials with binders and like known materials, followed by forming active material layer on collector. Binders are not specifically limited. Examples of binders are polytetrafluoroethylene, poly(vinylidene fluoride), and like fluorine-containing polymer; polyethylene, polypropylene, and like polyolefins; synthetic rubbers. The amount of binders are usually about 3-50 parts by

weight, preferably about 5-20 parts by weight, more preferably about 5-15 parts by weight based on 100 parts by weight of active material. An excessive amount of binder decreases a density of active material, thereby not preferable. A too small amount of binder has an insufficient ability to retain active material in electrode resulting in low stability of electrode, thereby not preferable. Examples of methods for producing an electrode are a method comprising producing paste by mixing active material and a binder, and forming an active material layer on collector with doctor blade or bar coater; and a method comprising adding a mixture of an active material and a binder to a press machine and forming a shaped form by pressing.

Known organic electrolyte solutions, inorganic solid electrolytes, solid polymer electrolytes may be used as electrolyte of lithium secondary battery of the invention.

The organic electrolyte solutions are in particular preferable from the viewpoint of ion conductivity. Examples of solvents for organic electrolyte solutions are propylene carbonate, ethylene carbonate, butylene carbonate, diethyl carbonate, dimethyl carbonate, methylethyl carbonate, γ -butyrolactone, and like esters; tetrahydrofuran, 2-

methyltetrahydrofuran and like substituted tetrahydrofuran; dioxolane, diethylether, dimethoxyethane, diethoxyethane, methoxyethoxyethane and like ethers; dimethylsulfoxide, sulfolane, methylsulfolane, acetonitrile, methyl formate and methyl acetate, which may be used singly or a mixture thereof. Examples of electrolytes are lithium perchlorate, lithium borofluoride, lithium hexafluorophosphate, lithium hexafluoroarsenate, sodium trifluoromethanesulfonate, lithium halide, lithium chloroalminate and like lithium salts, which are used singly or in a mixture thereof. Organic electrolyte solutions are prepared by dissolving electrolytes in said solvents. Solvents and electrolytes to prepare electrolyte solution is in no way limited to said examples.

Examples of inorganic solid electrolytes are nitride, halide, oxygen acid salts and phosphorous sulfide compounds of lithium, specifically, Li_3N , LiI , $\text{Li}_3\text{N-LiI-LiOH}$, LiSiO_4 , $\text{LiSiO}_4\text{-LiI-LiOH}$, $\text{Li}_3\text{PO}_4\text{-Li}_4\text{SiO}_4$ and Li_2SiS_3 .

Examples of organic solid electrolytes are substances comprising said electrolytes and dissociative polymers, and polymers with ion dissociative group or groups. The dissociative polymers include polyethyleneoxide derivatives and polymers including the

derivatives, polypropyleneoxide derivatives and polymers including the derivatives, poly(phosphate ester).

Polymer matrix material including said aprotic polar solvents, a mixture of polymer with ion dissociative group or groups and aprotic polar solvent, material prepared by adding polyacrylonitrile to electrolyte may be used. Inorganic solid electrolyte and organic solid electrolyte may be used in combination.

Positive electrode of lithium secondary battery of the invention may be prepared according to conventional manner using lithium-containing oxide as active material of positive electrode. Examples of active material of positive electrode are LiCoO_2 , LiNiO_2 , LiFeO_2 , LiMnO_2 , and analogs of the compounds $\text{Li}_x\text{M}_y\text{N}_z\text{O}_2$ wherein M represents any of Fe, Co, Ni and Mn, N represents a transition metal, 4B group metal or 5B group metal), LiVO_2 , which are mixed with electrically-conductive materials, binders and optionally with solid electrolytes to form a positive electrode. A mixing ratio of the materials is about 5-50 parts by weight of electrically-conductive materials, about 1-30 parts by weight of binder based on 100 parts by weight of active material. Such electrically-conductive materials are not specifically limited to, but include known carbon black (acetylene black, thermal black, channel black etc.) and

like carbons, graphite powder and metal powder. Binders are not specifically limited to, but include known polytetrafluoroethylene, poly(vinylidene fluoride), and like fluorine-containing polymers; polyethylene, polypropylene, and like polyolefins; and synthetic rubbers. In case that loadings of electrically-conductive material are less than 5 parts by weight or that loadings of binder are more than 30 parts by weight, practical lithium secondary battery can not be produced due to increased resistance or polarization of electrode and decreased discharge capacity. More than 50 parts by weight (relative ratio thereof is varied according to the type of electrically-conductive materials) of electrically-conductive material result in decreased amount of active material included in electrode leading to decreased discharge capacity of positive electrode. Less than 1 part by weight of binder results in insufficient integrity. More than 30 parts by weight of binder cause a decreased amount of active material included in electrode, an increased resistance or polarization of electrode and a decreased discharge capacity similar to electrically-conductive materials, thereby not practical. Production of positive electrode is conducted preferably by heat treatment near melting point of binder to improve integrity thereof.

Examples of separator to retain electrolyte solutions are known woven or unwoven fabric of electrical insulators such as synthetic resin fiber, glass fiber and natural fiber, powder molding of alumina and the like.

5 In particular, unwoven fabric of synthetic resin such as polyethylene and polypropylene are preferable from the viewpoint of stability of quality. An unwoven fabric of synthetic resin which is endowed with blocking function between positive and negative electrodes exerted by
10 melting the separator with abnormal exothermic heat of battery is suitably used from the viewpoint of safety. A thickness of separator is not specifically limited, as long as separator may retain a required amount of electrolyte solution and inhibit short circuit between
15 positive and negative electrodes, but generally about 0.01-1 mm, preferably about 0.02-0.05mm.

Materials of collector are not specifically limited to, but include known metals such as copper, nickel, stainless steel, aluminum and titanium in a form
20 of metallic foil, mesh, porous body and so on.

Effect of the Invention

According to the invention, novel carbon materials, in which surface of carbon materials to be core material is coated with heavy oil etc., may be
25 obtained by dipping carbon materials, in particular,

high-crystallinity graphite-type materials into tar, pitch and like coal heavy oil or petroleum heavy oil, followed by sepatrating coated carbon materials from heavy oil etc., washing with organic solvent and drying.

5 The carbon materials with a specific structure, wherein the surface of core material comprising graphite-type material with high crystallinity is covered by carbon-type material with low crystallinity, may be obtained by carbonization of graphite-type material whose
10 surface is uniformly coated with pitch at 600-2,000°C.

 According to the method for production of the invention, since no fusing or aggregation between particles occurs, when core carbon materials coated with pitch, tar and like heavy oil is washed, dried and
15 calcined, no grinding is required leading to preparation of near spheric, so called "edgeless", particles. The particles are free of deterioration factor, i.e., contamination during grinding.

 When non-aqueous secondary batteries or solid
20 electrolyte batteries are prepared using the coated carbon materials of the invention, in particular carbon materials prepared by coating surface of graphite material with heavy oil etc. or calcination product thereof, the batteries are superior in both charge and
25 discharge properties and safety.

The production method of the invention using inexpensive natural and artificial graphites and the like as core material, and inexpensive pitch, tar and the like as coating material is a simple and highly productive method so that inexpensive and high performance material for negative electrode of lithium secondary battery may be obtained.

Furthermore, according to the invention, possible are 4 combinations between core materials and surface materials including low-crystallinity carbon material + low-crystallinity carbon material; low-crystallinity carbon material + high-crystallinity carbon material; high-crystallinity carbon material and low-crystallinity carbon material; and high-crystallinity carbon material and high-crystallinity carbon material. Eight type of carbon materials are obtained in further consideration of 2 calcination step (carbonization step and graphitization step). Carbon materials comprising combinations of carbonization-treated high-crystallinity carbon material + low-crystallinity carbon material and high-crystallinity carbon material + high-crystallinity carbon material; and also graphitization-treated high-crystallinity carbon material + low-crystallinity carbon material are, in particular, useful as materials for negative electrode of lithium secondary battery because

of low reactivity with electrolyte and remarkable charge and discharge properties.

Best Mode for Carrying out the Invention

The inventions are described below in detail.

5 A variety of determinations are carried out according to the following ways.

1. Determination of particle diameter

Center particle diameter and particle size distribution of particles are determined with "FRA9220
10 Microtrac" product of Nikkiso.

2. Determination of covering ratio and coating thickness

With respect to carbon components from heavy oil covering the surface of core material before calcination, quinoline insoluble matter (%) was
15 determined by an analysis with solvent according to the method specified by JIS K2425 so as to calculate quinoline soluble matter (%) was calculated according to "100 - (quinoline insoluble matter)". The amount of quinoline soluble matter corresponds to the amount of
20 coat-forming carbon material.

A weight ratio of carbon materials for coating formation/(core carbon materials + carbon materials for coating formation) was calculated according to above-mentioned method.

25 3. Determination of specific surface area

The specific surface area was determined with "ASAP2400/nitrogen adsorption BET specific surface area measuring equipment" product of Micromeritics.

4. Determination of true specific gravity

5 True specific gravity was determined according to the method specified by JIS R7212.

5. Determination of crystal lattice size according to X-ray wide-angle diffractometry

10 Determination of crystal lattice size (L_c , L_a) according to X-ray wide-angle diffractometry was conducted by a known method described in "CARBON MATERIAL EXPERIMENTAL TECHNIQUE 1, pp.55-63, Ed. Carbon Material Society (KAGAKUGIYUTUSYA)". For the shape factor K , 0.90 was used.

15 6. Raman spectrometric analysis

Furthermore, as surface physical properties of carbon materials, R value was determined as ratio of peak strength of $1360\text{cm}^{-1}/1580\text{cm}^{-1}$ using 2 peaks observed by Raman spectrometric analysis with an argon laser at 20 514.5nm.

7. Determination of generated gas by immersing a negative electrode in electrolyte solution, followed by maintaining at elevated temperature

25 A pitch-coated carbon material (pitch-coated graphite) was calcined at $2,800\text{ }^{\circ}\text{C}$ for 1 hour under

nitrogen atmosphere for graphitization. 95 parts by weight of graphitized pitch-coated graphite and 5 parts by weight of dispersion-type PTFE ("D-1" product of Daikin Industries Ltd.) were mixed, uniformly stirred in liquid phase and dried to make a paste-like material. 0.25g of the material for negative electrode was molded by a pressing machine to produce a negative electrode body having a diameter of 2cm, and then dried in vacuo at 200°C for 6 hours.

Subsequently, the negative electrode was charged until electric potential thereof in electrolyte became 0 V. The charged negative electrode was placed into a beaker cell with 25 ml of electrolytic solution, which was heated at 60°C for 6 hours to determine the amount of generated gas per 1g of graphitized pitch-coated graphite.

As electrolytic solution, a mixed solvent of ethylene carbonate, diethyl carbonate and methyl propionate (3:3:4 ratio by volume) in which 1 mol dm^{-3} of LiClO_4 was dissolved was used.

8. Production of non-aqueous battery and determination of battery properties

A positive electrode is generally prepared by mixing material for positive electrode, electrically-conductive material and binder. Carbon black, graphite

and like carbon materials or metal powder, metal-wool,
and like metallic substances are suitably used as
electrically-conductive materials. Binders may be mixed
in the form of powder, or in the form of dispersion or
5 dissolved solution so as to improve integrity. When such
a dispersion or dissolved solution thereof is used,
removal of the solvent by vacuum treatment, heat
treatment or like means is required. Integrity may be
further improved by heat-treatment at near melting point
10 depending on the type of binders.

In examples of the present application, a
mixture of 100 parts by weight of LiCoO_2 as material for
positive electrode, 10 parts by weight of acetylene black
as electrically-conductive material and 10 parts by
15 weight of PTFE powder as binder was formed into electrode
having a diameter of 10mm to obtain a positive electrode
body.

A negative electrode body is produced as
follows in examples of the present application.

20 Pitch-coated graphite was calcined at $1,000^\circ\text{C}$
for 1 hour for carbonization. 95 parts by weight of the
carbonized pitch-coated graphite and 5 parts by weight of
dispersion-type PTFE ("D-1" product of Daikin Industries
Ltd.) were mixed, uniformly stirred in liquid phase and
25 dried to make paste-like material. 30mg of the material

for negative electrode was molded by a pressing machine to produce a negative electrode body having a diameter of 10mm, and then dried in vacuo at 200°C for 6 hours.

Separately, pitch-coated graphite was calcined at 2,800°C for 1 hour for graphitization. 95 parts by weight of the graphitized pitch-coated graphite and 5 parts by weight of dispersion-type PTFE ("D-1" product of Daikin Industries Ltd.) were mixed, uniformly stirred in liquid phase and dried to make paste-like material. 30mg of the material for negative electrode was molded by a pressing machine to produce a negative electrode body having a diameter of 10mm, and then dried in vacuo at 200°C for 6 hours.

Polypropylene unwoven fabric was used as separator.

Electrical discharge properties of coin-type lithium secondary battery, prepared by using the positive electrode body, the negative electrode body, the separator and the electrolytic solution thus obtained, were determined. The determination was carried out under constant-current charge and discharge of $1\text{mA}/\text{cm}^2$. Discharge capacity was regarded as capacity until battery voltage was decreased to 1.2V.

9. Preparation of solid electrolyte battery and determination of battery properties

The paste-like material of negative electrode prepared in the same manner as the item (said 8.) of preparation of non-aqueous battery was applied to both sides of copper foil in a thickness of 0.02 mm, dried and
5 rolled to obtain a negative electrode plate having a thickness of 0.10mm, width of 55mm and length of 90 mm.

A solid electrolyte, $(\text{PEO})_8 \cdot \text{LiClO}_4$, was prepared by dissolving polyethyleneoxide (molecular weight 600,000) and LiClO_4 in acetonitrile, followed by
10 casting the solution on a PTFE membrane ("TEFLON" product of DUPONT) under argon atmosphere in glovebox, and then distillating the solvent by allowing it to stand at 25 °C in glovebox.

A film-type lithium secondary battery was
15 prepared by using carbonized pitch-coated graphite or graphitized pitch-coated graphite as negative electrode body, LiCoO_2 as solid electrolyte and positive electrode body and $(\text{PEO})_8 \cdot \text{LiClO}_4$ as solid electrolyte.

Electrical discharge properties of
20 lithium secondary battery thus prepared were determined. The determination was carried out under constant-current charge and discharge of $1\text{mA}/\text{cm}^2$. Discharge capacity was regarded as capacity until battery voltage was decreased to 1.2V.

25 Example 1

50g of massive artifitial graphite (center
particle diameter D50 = 7.5 μ m; particle size distribution
0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific
surface area = 10.8m²/g; R value = 0.26; true specific
5 gravity = 2.25g/cm²) and 100g of coal tar pitch
(softening point 80°C; quinoline insoluble matter =
trace; toluene insoluble matter = 30%) whose primary QI
was removed previously were added to a 500ml reaction
flask. The mixture was stirred for 2 hours at 200°C
10 under normal pressure to give crude pitch-coated
graphite. Toluene (1 part) was added to the crude pitch-
coated graphite thus obtained (1 part), and the mixture
was stirred at 80°C for 1 hour for washing treatment and
filtered to give purified pitch-coated graphite. Center
15 particle diameter D50 of the purified pitch-coated
graphite was determined as 7.7 μ m. Since center particle
diameter D50 of graphite as core material was 7.5 μ m, a
thickness of pitch layer is 0.1 μ m.

The quinoline soluble matter, specific surface
20 area and true specific gravity of the resulting purified
pitch-coated graphite are shown in table 1. Since
quinoline soluble matter was 9.6%, a covering ratio of
the purified pitch-coated graphite is 0.096.

The purified pitch-coated graphite was calcined
25 at 1,000°C for 1 hour under nitrogen atmosphere (heating

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rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1µm or less of the carbonized pitch-coated graphite are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite has a distribution within 0.1-150 µm. The results of X-ray diffractometry thereof were similar to those of core material.

Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicate that artificial graphite as core material was coated with carbonized pitch forming a coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 mol dm^{-3} of LiClO_4 was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Furthermore, a negative electrode was produced using a negative electrode which was produced by using

the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 3.

Example 2

5 The purified pitch-coated graphite obtained according to example 1 was calcined at 1,000°C for 1 hour under vacuum of 10 torr (heating rate 25°C/hr) for vacuum carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of
10 particles having a diameter of 1 μ m or less of the resulting vacuum carbonized pitch-coated graphite are shown in table 1. The results of determination of particle size distribution of the vacuum carbonized pitch-coated graphite indicated that the coated graphite
15 had a distribution within 0.1-150 μ m as core material. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between core material and vacuum carbonized pitch-coated graphite indicated that carbonized pitch forming a
20 coating layer had lower crystallinity than core material. The results of SEM observation indicated that artificial graphite as core material are coated with vacuum carbonized pitch forming a coating layer and that edge parts thereof were rounded.

25 A non-aqueous secondary battery was produced

using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the vacuum carbonized pitch-coated graphite. The results of
5 determination of charge and discharge properties thereof are shown in table 2.

Example 3

The purified pitch-coated graphite obtained according to example 1 was calcined at 2,800°C for 1 hour
10 under nitrogen atmosphere for graphitization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1µm or less of the resulting graphatized pitch-coated graphite are shown in table 1. The results
15 of determination of particle size distribution of the graphatized pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 µm as core material. The results of X-ray diffractometry thereof were similar to those of core material.
20 Furthermore, comparison of R values between core material and graphatized pitch-coated graphite indicated that graphitized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core
25 material was coated with graphitized pitch forming a

coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, a mixed solvent of ethylene carbonate, diethyl carbonate and methyl-
5 propionate (3:3:4) in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the graphatized pitch-coated graphite.

The amount of generated gas of the graphatized pitch-coated graphite in the electrolytic solution was
10 determined. The results of determination of charge and discharge properties and the amount of generated gas are shown in table 2.

Example 4

The purified pitch-coated graphite obtained
15 according to example 1 was calcined at 1,000°C in a lead furnace capable of elevating temperature very slowly (reducing atmosphere, heating rate of up to 5°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of
20 particles having a diameter of 1μm or less of the resulting carbonized pitch-coated graphite are shown in table 1. The results of determination of particle size distribution of the carbonized pitch-coated graphite indicated that the coated graphite had a distribution
25 within 0.1-150 μm as core material. The results of X-ray

diffractionmetry thereof were similar to those of core material. Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch forming a coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 5

The purified pitch-coated graphite obtained according to example 1 was calcined at 1,300°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1µm or less of the carbonized pitch-coated graphite are shown in table 1. The results of determination of particle size distribution of the carbonized pitch-coated graphite

indicated that the coated graphite had a distribution within 0.1-150 μm as core material. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between

5 core material and carbonized pitch-coated graphite indicated that carbonized pitch forming a coating layer has lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch forming a
10 coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO_4 was dissolved and a negative electrode which was produced by using the carbonized
15 pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 6

The purified pitch-coated graphite obtained
20 according to example 1 was treated for oxidation at 300°C for 8 hours under air atmosphere in a chamber with constant temperature and relative humidity. The covering ratio, specific surface area and true specific gravity of the resulting oxidized and purified pitch-coated graphite
25 are shown in table 1. The oxidized and purified pitch-

coated graphite was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1µm or less of the carbonized pitch-coated graphite are shown in table 1. The results of determination of particle size distribution of the carbonized pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 µm as core material. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch forming a coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 7

50g of massive artificial graphite (center particle diameter D50 = 7.5 μ m; particle size distribution 0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³) and 100g of coal tar pitch (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 500ml reaction flask. The mixture was stirred for 2 hours at 200°C to give crude pitch-coated graphite.

Toluene (1 part) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 20°C for 1 hour for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as 7.9 μ m. Since center particle diameter D50 of graphite as core material was 7.5 μ m, a thickness of pitch layer is 0.2 μ m.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline soluble matter thereof was 20.4%, a covering ratio of the purified pitch-coated graphite is 0.204.

The purified pitch-coated graphite thus

obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less with respect to the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μ m. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch forming a coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 8

The purified pitch-coated graphite obtained according to example 7 was calcined at 2,800°C for 1 hour under nitrogen atmosphere for graphitization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1µm or less of the resulting graphitized pitch-coated graphite are shown in table 1. The results of determination of particle size distribution of the graphitized pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 µm as core material. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between core material and graphitized pitch-coated graphite indicated that graphitized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artificial graphite as core material was coated with graphitized pitch forming a coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, a mixed solvent of ethylene carbonate, diethyl carbonate and methylpropionate (3:3:4) in which 1 mol/dm³ of LiClO₄ was dissolved and a negative electrode which was produced by

using the graphatized pitch-coated graphite. The amount of generated gas of the graphatized pitch-coated graphite in the electrolytic solution was determined. The results of determination of charge and discharge properties and the amount of generated gas are shown in table 2.

Example 9

50g of massive artifitial graphite (center particle diameter D50 = 7.5 μ m; particle size distribution 0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³) and 100g of coal tar pitch (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 500ml reaction flask. The mixture was stirred for 2 hours at 200°C under reduced pressure (evacuated with a vacuum pump; reduced pressure = 50 torr) to give crude pitch-coated graphite.

Toluene (1 part) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 80°C for 1 hour for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as 7.7 μ m. Since center particle diameter D50 of graphite as core material was 7.5 μ m, a

thickness of pitch layer is 0.1 μ m.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline
5 soluble matter thereof was 10.4%, a covering ratio of the purified pitch-coated graphite is 0.104.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for
10 carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less of the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size
15 distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μ m as core material. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between
20 core material and carbonized pitch-coated graphite indicated that carbonized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch forming a
25 coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 10

50g of massive artificial graphite (center particle diameter D50 = 7.5μm; particle size distribution 0.1-150μm; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm²) and 100g of coal tar pitch (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 500ml reaction flask. The mixture was stirred for 2 hours at 200°C to give crude pitch-coated graphite.

Tar middle oil (1 part) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 20°C for 1 hour for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as 7.6μm. Since center particle diameter D50 of graphite as core material

is 7.5 μ m, a thickness of pitch layer is 0.05 μ m.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline soluble matter thereof was 8.8%, a covering ratio of the purified pitch-coated graphite is 0.088.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less of the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μ m as core material. The results of X-ray diffractometry thereof are similar to those of core material. Furthermore, comparison of R values between core material and graphitized pitch-coated graphite indicated that graphitized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artificial graphite as core material was coated with graphitized pitch forming a coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 11

50g of massive artificial graphite (center particle diameter D50 = 7.5μm; particle size distribution 0.1-150μm; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³) and 200g of coal tar pitch (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 1,000ml reaction flask. The mixture was stirred for 2 hours at 200°C to give crude pitch-coated graphite.

Toluene (1 part) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 80°C for 1 hour for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as 7.9μm. Since center particle diameter D50 of graphite as core material is 7.5μm,

thickness of pitch layer is 0.2 μ m.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline
5 soluble matter thereof was 17.3%, a covering ratio of the purified pitch-coated graphite is 0.173.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for
10 carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less of the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size
15 distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μ m as core material. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between
20 core material and graphitized pitch-coated graphite indicated that carbonized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch forming a
25 coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 12

50g of massive artificial graphite (center particle diameter D50 = 7.5μm; particle size distribution 0.1-150μm; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³) and 100g of coal tar pitch (softening point 80°C; quinoline insoluble matter = 3.9%; toluene insoluble matter = 34%) whose primary QI was not removed were added to a 500ml reaction flask. The mixture was stirred for 2 hours at 200°C under normal pressure to give crude pitch-coated graphite.

Toluene (1 part) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 80°C for 1 hour for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as 7.9μm. Since center particle diameter D50 of graphite as core material was 7.5μm,

thickness of pitch layer is 0.2 μ m.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline
5 soluble matter thereof was 7.5%, a covering ratio of the purified pitch-coated graphite is 0.075.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for
10 carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less with respect to the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of
15 particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μ m as core material. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R
20 values between core material and carbonized pitch-coated graphite indicated that carbonized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artificial graphite as core material was coated with carbonized
25 pitch forming a coating layer and that edge parts thereof

were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 13

50g of massive artificial graphite (center particle diameter D50 = 7.5μm; particle size distribution 0.1-150μm; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm²) and 100g of coal tar pitch (softening point 10°C; quinoline insoluble matter = trace; toluene insoluble matter = 8%) whose primary QI was removed previously were added to a 500ml reaction flask. The mixture was stirred for 2 hours at 200°C under normal pressure to give crude pitch-coated graphite.

Toluene (1 part) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 80°C for 1 hour for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated

graphite was determined as 7.6 μ m. Since center particle diameter D50 of graphite as core material is 7.5 μ m, a thickness of pitch layer is 0.05 μ m.

5 The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline soluble matter thereof was 7.8%, a covering ratio of the purified pitch-coated graphite is 0.078.

10 The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less of the
15 carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μ m as core material. The results of X-ray
20 diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch forming a coating layer had lower crystallinity than core material. The results
25 of SEM observation indicated that artifitial graphite as

core material was coated with carbonized pitch forming a coating layer and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Furthermore, a negative electrode which was produced by using the carbonized pitch-coated graphite, and then a solid electrolyte lithium secondary battery was produced. The results of determination of charge and discharge properties thereof are shown in table 3.

Example 14

50g of graphitized spheric mesocarbon microbeads ("MCMB-6-28" product of Osaka Gas Co., Ltd., center particle diameter D50 = 6.0μm; particle size distribution 0.1-50μm; d002=0.336nm; Lc=50nm; La=90nm; specific surface area = 3.0m²/g; R value = 0.42; true specific gravity = 2.20g/cm³) and 100g of coal tar pitch (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 500ml reaction flask. The mixture was stirred for 2 hours at 200°C

under normal pressure to give crude pitch-coated mesocarbon microbeads.

Toluene (1 part) was added to the crude pitch-coated mesocarbon microbeads thus obtained (1 part), and the mixture was stirred at 80°C for 1 hour for washing treatment and filtered to give purified pitch-coated mesocarbon microbeads. Center particle diameter D50 of the purified pitch-coated mesocarbon microbeads was determined as 6.2µm. Since center particle diameter D50 of graphite as core material was 6.0µm, a thickness of pitch layer is 0.1µm.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated mesocarbon microbeads obtained are shown in table 1. Since quinoline soluble matter thereof was 9.8%, a covering ratio thereof is 0.098.

The purified pitch-coated graphitized mesocarbon microbeads thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1µm or less of the purified pitch-coated graphitized mesocarbon microbeads are shown in table 1. The results of determination of particle size distribution of the

purified pitch-coated graphitized mesocarbon microbeads indicated that the mesocarbon microbeads had a distribution within 0.1-50 μm as core material.

Furthermore, comparison of R values between core material

5 and carbonized pitch-coated graphitized mesocarbon microbeads indicated that carbonized pitch forming a coating layer had lower crystallinity than core material.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate
10 in which 1 mol dm^{-3} of LiClO_4 was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphitized mesocarbon microbeads. The results of determination of charge and discharge properties thereof are shown in table 2.

15 Example 15

50g of massive artificial graphite (center particle diameter D50 = 16.2 μm ; particle size distribution 0.1-120 μm ; d002=0.337nm; Lc=100nm; La=71nm; specific surface area = 14.4 m^2/g ; R value = 0.31; true
20 specific gravity = 1.96 g/cm^3) and 100g of coal tar pitch (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 1,000ml reaction flask. The mixture was stirred for 5 hours at 250°C
25 under normal pressure to give crude pitch-coated

graphite.

Toluene (3 parts) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 50°C for 5 hour for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as 16.6 μ m. Since center particle diameter D50 of graphite as core material is 16.2 μ m, a thickness of pitch layer is 0.2 μ m.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline soluble matter thereof was 11.3%, a covering ratio of the purified pitch-coated graphite is 0.113.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less with respect to the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-120 μ m as core material. The

results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch forming a coating layer had lower crystallinity than core material. The results of SEM observation indicated that artificial graphite as core material was coated with carbonized pitch, i.e., coat-forming carbon materials and that edge parts thereof were rounded.

10 A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 16

50g of massive artificial graphite (center particle diameter D50 = 16.2μm; particle size distribution 1-80μm; d002=0.338nm; Lc=83nm; La=63nm; specific surface area = 6.8m²/g; R value = 0.38; true specific gravity = 2.02g/cm³) and 100g of coal tar pitch (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 1,000ml reaction

flask. The mixture was stirred for 5 hours at 250°C under normal pressure to give crude pitch-coated graphite.

Toluene (3 parts) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 50°C for 5 hour for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as 12.0µm. Since center particle diameter D50 of graphite as core material was 11.6µm, a thickness of pitch layer is 0.2µm.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline soluble matter thereof was 12.3%, a covering ratio of the purified pitch-coated graphite is 0.123.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1µm or less with respect to the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated

graphite indicated that the coated graphite had a distribution within 1-80 μm . The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between
5 core material and carbonized pitch-coated graphite indicated that carbonized pitch, i.e., coat-forming carbon materials had lower crystallinity than core material. The results of SEM observation indicated that
10 artificial graphite as core material was coated with carbonized pitch, i.e., coat-forming carbon materials and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 mol dm^{-3} of LiClO_4 was dissolved and a negative
15 electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 17

20 50g of scaly artificial graphite (center particle diameter D50 = 18.9 μm ; particle size distribution 0.1-150 μm ; d002=0.340nm; Lc=42nm; La=50nm; specific surface area = 9.2 m^2/g ; R value = 0.49; true specific gravity = 1.82 g/cm^3) and 100g of coal tar pitch
25 (softening point 80°C; quinoline insoluble matter =

trace; toluene insoluble matter = 30%) whose primary QI
was removed previously were added to a 1,000ml reaction
flask. The mixture was stirred for 5 hours at 250°C
under normal pressure to give crude pitch-coated
5 graphite.

Toluene (3 parts) was added to the crude pitch-
coated graphite thus obtained (1 part), and the mixture
was stirred at 50°C for 5 hour for washing treatment and
filtered to give purified pitch-coated graphite. Center
10 particle diameter D50 of the purified pitch-coated
graphite was determined as 19.3µm. Since center particle
diameter D50 of graphite as core material was 18.9µm, a
thickness of pitch layer is 0.2µm.

The covering ratio, specific surface area and
15 true specific gravity of the purified pitch-coated
graphite obtained are shown in table 1. Since quinoline
soluble matter thereof was 10.6%, a covering ratio of the
purified pitch-coated graphite is 0.106.

The purified pitch-coated graphite thus
20 obtained was calcined at 1,000°C for 1 hour under
nitrogen atmosphere (heating rate 25°C/hr) for
carbonization. The specific surface area, true specific
gravity, R value and volume-based integrated value of
particles having a diameter of 1µm or less of the
25 carbonized pitch-coated graphite obtained are shown in

table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μm . The results of X-ray diffractometry thereof were similar to those of core material.

Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch, i.e., coat-forming carbon materials had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch, i.e., coat-forming carbon materials and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 mol dm^{-3} of LiClO_4 was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 18

50g of whisker artifitial graphite (center particle diameter D50 = 23.8 μm ; particle size distribution 0.1-150 μm ; d_{002} =0.347nm; L_c =25nm; L_a =15nm; specific surface area = 13.5 m^2/g ; R value = 0.68; true

specific gravity = 1.60g/cm^3) and 100g of coal tar pitch (softening point 80°C ; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 1,000ml reaction
5 flask. The mixture was stirred for 5 hours at 250°C under normal pressure to give crude pitch-coated graphite.

Toluene (3 parts) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture
10 was stirred at 50°C for 5 hours for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as $24.2\mu\text{m}$. Since center particle diameter D50 of graphite as core material was $23.8\mu\text{m}$, a
15 thickness of pitch layer is $0.2\mu\text{m}$.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline soluble matter thereof was 13.1%, a covering ratio of the
20 purified pitch-coated graphite is 0.131.

The purified pitch-coated graphite thus obtained was calcined at $1,000^\circ\text{C}$ for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific
25 gravity, R value and volume-based integrated value of

particles having a diameter of $1\mu\text{m}$ or less of the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite

5 indicated that the coated graphite had a distribution within $0.1\text{--}150\mu\text{m}$. The results of X-ray diffractometry thereof were similar to those of core material.

Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that
10 carbonized pitch, i.e., coat-forming carbon materials had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch, i.e., coat-forming carbon materials and that edge parts thereof
15 were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm^{-3} of LiClO_4 was dissolved and a negative electrode which was produced by using the carbonized
20 pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 19

50g of massive artifitial graphite (center
25 particle diameter $D_{50} = 7.5\mu\text{m}$; particle size distribution

0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³) and 100g of coal tar pitch

(softening point 80°C; quinoline insoluble matter =

5 trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 500ml reaction flask. The mixture was stirred for 1 hour at 300°C under normal pressure to give crude pitch-coated graphite.

Quinoline (0.1 part) was added to the crude
10 pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 150°C for 10 hours for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified
15 pitch-coated graphite was determined as 8.1 μ m. Since center particle diameter D50 of graphite as core material was 7.5 μ m, thickness of pitch layer is 0.3 μ m.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline
20 soluble matter thereof was 29.0%, a covering ratio of the purified pitch-coated graphite is 0.290.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 100°C/hr) for
25 carbonization. The specific surface area, true specific

gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less of the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μ m. The results of X-ray diffractometry thereof were similar to those of core material.

Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch, i.e., coat-forming carbon materials had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch, i.e., coat-forming carbon materials and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 20

25g of massive artifitial graphite (center

particle diameter D50 = 7.5 μ m; particle size distribution 0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³) and 50g of coal tar pitch (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 1,000ml reaction flask. The mixture was stirred for 3 hours at 30°C under normal pressure to give crude pitch-coated graphite.

10 Acetone (10 parts) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 30°C for 5 hours for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified
15 pitch-coated graphite was determined as 7.8 μ m. Since center particle diameter D50 of graphite as core material was 7.5 μ m, a thickness of pitch layer is 0.15 μ m.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated
20 graphite obtained are shown in table 1. Since quinoline soluble matter thereof was 15.0%, a covering ratio of the purified pitch-coated graphite is 0.150.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under
25 nitrogen atmosphere (heating rate 25°C/hr) for

carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less of to the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μ m. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch, i.e., coat-forming carbon materials had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch, i.e., coat-forming carbon materials and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 21

50g of massive artifitial graphite (center particle diameter D50 = 7.5 μ m; particle size distribution 0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³) and 50g of coal tar pitch (softening point 10°C; quinoline insoluble matter = trace; toluene insoluble matter = 8%) whose primary QI was removed previously were added to a 500ml reaction flask. The mixture was stirred for 3 hours at 250°C under normal pressure to give crude pitch-coated graphite.

Tar middle oil (10 parts) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 200°C for 1 hour for washing treatment and filtered to give purified pitch-coated graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as 7.5 μ m.

The covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline soluble matter thereof was 2.0%, a covering ratio of the purified pitch-coated graphite is 0.020.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific

gravity, R value and volume-based integrated value of particles having a diameter of 1 μ m or less of the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 μ m. The results of X-ray diffractometry thereof were similar to those of core material.

Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch, i.e., coat-forming carbon materials had lower crystallinity than core material. The results of SEM observation indicated that artifitial graphite as core material was coated with carbonized pitch, i.e., coat-forming carbon materials and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 22

50g of massive artifitial graphite (center

particle diameter D50 = 7.5 μ m; particle size distribution 0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³) and 100g of coal tar pitch

5 (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 8%) whose primary QI was removed previously were added to a 1,000ml reaction flask. The mixture was stirred for 3 hours at 250°C under normal pressure to give crude pitch-coated
10 graphite.

Toluene (4 parts) was added to the crude pitch-coated graphite thus obtained (1 part), and the mixture was stirred at 80°C for 1 hour for washing treatment and filtered to give purified pitch-coated
15 graphite. Center particle diameter D50 of the purified pitch-coated graphite was determined as 7.6 μ m. Since center particle diameter D50 of graphite as core material was 7.5 μ m, a thickness of pitch layer is 0.05 μ m.

The covering ratio, specific surface area and
20 true specific gravity of the purified pitch-coated graphite obtained are shown in table 1. Since quinoline soluble matter thereof was 8.2%, a covering ratio of the purified pitch-coated graphite is 0.082.

The purified pitch-coated graphite thus
25 obtained was calcined at 700°C for 1 hour under nitrogen

atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1µm or less of the carbonized pitch-coated graphite obtained are shown in table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 µm. The results of X-ray diffractometry thereof were similar to those of core material. Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch, i.e., coat-forming carbon materials had lower crystallinity than core material. The results of SEM observation indicated that artificial graphite as core material was coated with carbonized pitch, i.e., coat-forming carbon materials and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 mol dm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

Example 23

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized
25 pitch-coated graphite. The results of determination of

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using the carbonized
25 pitch-coated graphite. The results of determination of

charge and discharge properties thereof are shown in table 2.

Example 24

50g of massive artificial graphite (center
particle diameter D50 = 7.5 μ m; particle size distribution
0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific
surface area = 10.8m²/g; R value = 0.26; true specific
gravity = 2.25g/cm³) and 100g of coal tar pitch
(softening point 10°C; quinoline insoluble matter = 2.9%;
toluene insoluble matter = 7.8%) whose primary QI was
adjusted previously were added to a 1,000ml reaction
flask. The mixture was stirred for 2 hours at 200°C
under normal pressure to give crude pitch-coated
graphite.

Toluene (4 parts) was added to the crude
pitch-coated graphite thus obtained (1 part), and the
mixture was stirred at 80°C for 1 hour for washing
treatment and filtered to give purified pitch-coated
graphite. Center particle diameter D50 of the purified
pitch-coated graphite was determined as 7.6 μ m. Since
center particle diameter D50 of graphite as core material
was 7.5 μ m, a thickness of pitch layer is 0.05 μ m.

The covering ratio, specific surface area and
true specific gravity of the purified pitch-coated
graphite obtained are shown in table 1. Since quinoline

soluble matter thereof was 8.7%, a covering ratio of the purified pitch-coated graphite is 0.087.

The purified pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under
5 nitrogen atmosphere (heating rate 25°C/hr) for carbonization. The specific surface area, true specific gravity, R value and volume-based integrated value of particles having a diameter of 1µm or less of the carbonized pitch-coated graphite obtained are shown in
10 table 1. The results of determination of particle size distribution of the purified pitch-coated graphite indicated that the coated graphite had a distribution within 0.1-150 µm. The results of X-ray diffractometry thereof were similar to those of core material.
15 Furthermore, comparison of R values between core material and carbonized pitch-coated graphite indicated that carbonized pitch, i.e., coat-forming carbon materials had lower crystallinity than core material. The results of SEM observation indicated that artificial graphite as
20 core material was coated with carbonized pitch, i.e., coat-forming carbon materials and that edge parts thereof were rounded.

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate
25 in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative

electrode which was produced by using the carbonized pitch-coated graphite. The results of determination of charge and discharge properties thereof are shown in table 2.

5 Comparative example 1

A non-aqueous secondary battery was produced using, as an electrolytic solution, propylene carbonate in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using massive artifitial graphite (center particle diameter D50 = 7.5μm; particle size distribution 0.1-150μm; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³).

However, the battery can hardly be charged and discharged due to decomposition of the electrolytic solution.

A covering ratio, specific surface area and true specific gravity of the purified pitch-coated graphite obtained are shown in table 1.

20 Comparative example 2

A non-aqueous secondary battery was produced using, as an electrolytic solution, a mixed solvent of ethylene carbonate, diethyl carbonate and methylpropionate (3:3:4) in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by

using massive artifitial graphite (center particle diameter D50 = 7.5 μ m; particle size distribution 0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³). The amount of generated gas of the graphite in the electrolytic solution was determined. The results of determination of charge and discharge properties and the amount of generated gas are shown in table 2.

Comparative example 3

A negative electrode was produced by using massive artifitial graphite (center particle diameter D50 = 7.5 μ m; particle size distribution 0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³). A solid electrolyte lithium secondary battery was then produced using the negative electrode. The results of determination of charge and discharge properties thereof are shown in table 3.

Comparative example 4

A non-aqueous secondary battery was produced using, as an electrolytic solution, a mixed solvent of ethylene carbonate, diethyl carbonate and methylpropionate (3:3:4) in which 1 moldm⁻³ of LiClO₄ was dissolved and a negative electrode which was produced by using graphitized spheric mesocarbon microbeads ("MCMB-6-

28" product of Osaka Gas Co., Ltd., center particle diameter D50 = 6.0 μ m; particle size distribution 0.1-50 μ m; d002=0.336nm; Lc=50nm; La=90nm; specific surface area = 3.0m²/g; R value = 0.42; true specific gravity = 2.20g/cm³). The results of determination of charge and discharge properties thereof are shown in table 2.

Comparative example 5

50g of massive artificial graphite (center particle diameter D50 = 7.5 μ m; particle size distribution 0.1-150 μ m; d002=0.336nm; Lc=100nm; La=97nm; specific surface area = 10.8m²/g; R value = 0.26; true specific gravity = 2.25g/cm³) and 100g of coal tar pitch (softening point 80°C; quinoline insoluble matter = trace; toluene insoluble matter = 30%) whose primary QI was removed previously were added to a 1,000ml reaction flask. The mixture was stirred for 2 hours at 200°C under normal pressure to give crude pitch-coated graphite.

The crude pitch-coated graphite thus obtained was calcined at 1,000°C for 1 hour under nitrogen atmosphere (heating rate 25°C/hr) for carbonization without washing the crude pitch-coated graphite with organic solvents. After calcination, the artificial graphite powder was taken out as lump. The lump of carbon material was grinded with a coffee mill to obtain

carbon material powders. The specific surface area, true
specific gravity, R value and volume-based integrated
value of particles having a diameter of $1\mu\text{m}$ or less of
the obtained carbon materials are shown in table 1. The
5 results indicate small R value thereof. The SEM
observation demonstrates that the carbon materials has an
angular shape in the carbon materials produced according
to the method of the invention. The angular shape seems
to result from exposure of graphite face caused by
10 grinding.

A non-aqueous secondary battery was produced
using, as an electrolytic solution, propylene carbonate
in which 1 mol dm^{-3} of LiClO_4 was dissolved and a negative
electrode which was produced by using the carbon
15 materials. The results of determination of charge and
discharge properties thereof are shown in table 2.

Table 1

	cover- ing ratio	Parti- cle diam- eter	Parti- cle less than 1µm (vol%)	R value	Specific surface area		True specific gravity	
					Before calci- nation	After calci- nation	Before calci- nation	After calci- nation
5								
	Ex.1	Ex.2	Ex.3	Ex.4	Ex.5	Ex.6	Ex.7	Ex.8
	Ex.9	Ex.10	Ex.11	Ex.12	Ex.13	Ex.14	Ex.15	Ex.16
10	Ex.17	Ex.18	Ex.19	Ex.20	Ex.21	Ex.22	Ex.23	Ex.24
	Comp.Ex.1	Comp.Ex.1	Comp.Ex.1	Comp.Ex.3	Comp.Ex.4	Comp.Ex.4	Comp.Ex.5	
	trace	trace	trace	trace	trace			

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Table 2

		Discharge capacity (mAh/g)	Efficiency of charge and discharge(%)	Generated gas (ml/g)
5	Example 1	355	83.4	-
	Example 2	370	86.0	-
	Example 3	343	86.2	2.2
	Example 4	346	85.9	-
	Example 5	349	86.1	-
10	Example 6	359	85.3	-
	Example 7	342	85.5	-
	Example 8	334	90.5	1.7
	Example 9	348	87.8	-
	Example 10	351	84.2	-
15	Example 11	344	89.5	-
	Example 12	339	82.4	-
	Example 13	342	90.1	-
	Example 14	334	90.5	-
	Example 15	330	82.5	-
20	Example 16	316	82.1	-
	Example 17	310	87.6	-
	Example 18	303	88.7	-
	Example 19	331	88.6	-
	Example 20	340	82.7	-
25	Example 21	357	80.6	-
	Example 22	321	83.0	-
	Example 23	359	80.4	-
	Example 24	350	82.8	-
	Comp.Ex.1	Determination is impossible due to decomposition of solvent		
30	Comp.Ex.2	224	51.0	15.0
	Comp.Ex.4	303	90.2	-
	Comp.Ex.5	302	68.1	-

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Table 3

	Discharge capacity (mAh/g)	Efficiency of Charge and Discharge(%)
Ex. 1	350	83.1
5 Comp.ex. 3	210	49.0

As shown in table 1, coating a graphite surface by pitch or tar can reduce specific surface area thereof. The specific surface area is further reduced by calcination of the coated graphite.

10 As shown in table 2, coating a graphite surface with pitch or tar greatly improves discharge capacity and efficiency of charge and discharge of non-aqueous lithium secondary battery. Coating a graphite surface by pitch inhibits a reactivity of the graphite to the electrolytic
15 solution and reduces the amount of generated gas.

As shown in table 3, discharge capacity and efficiency of charge and discharge of battery may be greatly improved by coating a graphite surface with pitch or tar with respect to solid electrolyte lithium
20 secondary battery.

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CLAIMS (Art. 34 amendment (2) filed on Jan. 12, 1998)

1. A carbon material characterized in that
edge parts of a core carbon material are partially or
entirely coated with coat-forming carbon material and
5 that the carbon material is nearly spheric or
ellipsoidal.

2. The carbon material according to claim 1
wherein the carbon material has a specific surface area
determined by a BET method of $5\text{m}^2/\text{g}$ or less.

10 3. The carbon material according to claim 1 or
2 wherein the coat-forming carbon material has a lower
crystallinity than the core carbon material.

4. The carbon material according to any of
claims 1-3 wherein the core carbon material is a carbon
15 material with high crystallinity having a mean
interplanar spacing (d002) of (002) plane of 0.335-0.340
nm, a thickness of crystallite size in direction of (002)
plane (Lc) of at least 10nm, a thickness of crystallite
size in direction of (110) plane (La) of at least 10 nm.

20 5. The carbon material according to any of
claims 1-4 wherein a true specific gravity of entire
carbon material is 1.50 to $2.26\text{g}/\text{cm}^3$.

6. The carbon material according to any of
claims 1-5 wherein a volume-based integrated value of
25 particles having a diameter of $1\mu\text{m}$ or less determined by

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12. The calcined two-layer carbon material according to any of claims 7-11 wherein a volume-based integrated value of particles having a diameter of 1 μ m or less determined by particle size distribution is 10% or less.

13. A method for producing a carbon material having a coating layer on the surface characterized in that the method comprises dipping a core carbon material into a coat-forming carbon material, separating the core carbon material from the coat-forming carbon material, adding organic solvent or solvents to the separated core carbon material which is subjected to washing, drying and calcination.

14. A method for producing a carbon material
15 having a coating layer on the surface characterized in
that the method comprises dipping a core carbon material
into a coat-forming carbon material at 10-300°C,
separating the core carbon material from the coat-forming
carbon material, adding organic solvent or solvents to
20 the separated core carbon material which is subjected to
washing, drying and calcination.

15. The method for producing a coated carbon material according to claim 13 or 14, wherein the separated core carbon material to which the organic solvent or solvents are added is washed at 10-300°C.

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16. The method for producing a coated carbon material according to any of claims 13-15, wherein the core carbon material is dipped into the coat-forming carbon material under reduced pressure.

5 17. The method for producing a coated carbon material according to any of claims 13-16, wherein the coat-forming carbon material is coal heavy oil or petroleum heavy oil.

10 18. The method for producing a coated carbon material according to any of claims 13-17, wherein the coat-forming carbon material is tar or pitch.

15 19. The method for producing a coated carbon material according to any of claims 13-18, wherein the organic solvents used for washing are at least one selected from toluene, quinoline, acetone, hexane, benzene, xylene, methylnaphthalene, alcohols, oils from coal and petroleum.

20 20. The method for producing a coated carbon material according to any of claims 13-19, wherein a ratio of solid matter and organic solvent or solvents during washing is 1:0.1-10 by weight.

25 21. The method for producing a coated carbon material according to any of claims 13-20, wherein a covering ratio (c) defined as weight ratio of coat-forming carbon material/(core carbon material +

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27. A method for producing a coated carbon material according to any of claims 23-26, wherein a surface of the coated carbon material is pretreated for

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5 29. A lithium secondary battery comprising
material for negative electrode comprising the carbon
material according to any of claims 1-12.

31. The lithium secondary battery according to claim 30, wherein the lithium secondary battery is a solid electrolyte lithium secondary battery.

33. A method for producing a two-layer carbon material characterized in that the coated carbon material produced according to claim 32 is calcined for

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5 carbonization at a heating rate of up to 10°C/hr.

10 36. A method for producing a two-layer carbon material characterized in that the coated carbon material produced according to claims 32 is calcined for graphitization.

37. The method for producing a two-layer
15 carbon material according to any of claims 32-36, wherein
a surface of the coated carbon material is pretreated for
oxidation before calcination of the coated carbon
material.

DECLARATION FOR USA PATENT APPLICATION

(including Design and National Stage PCT)

Attorney's Docket ID: _____

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below adjacent to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention

entitled MATERIAL FOR NEGATIVE ELECTRODE OF LITHIUM SECONDARY BATTERY, METHOD FOR PRODUCTION THEREOF AND LITHIUM SECONDARY BATTERY USING THE SAME, the specification of which

(check one) ☐ is attached hereto.

☐ is U.S. Application Serial No. _____ which was filed on _____ and (if applicable) amended on _____.

☒ is International Application No. PCT/JP96/03344 which was filed on November 14, 1996 and (if applicable) amended on Apr. 24, 1997, Nov. 17, 1997 and Jan. 12, 1998.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, § 1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, § 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Prior Foreign Application(s) (____ ADDITIONAL APPLICATIONS IDENTIFIED ON ATTACHED SHEET):

Number	Country	Day/Month/Year Filed	Priority Claimed Yes No
295462/1995	Japan	14/11/1995	X

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) listed below; and insofar as the subject matter of each claim of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations § 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application in accordance with 37 C.F.R. 1.63(d). (____ ADDITIONAL APPLICATIONS IDENTIFIED ON ATTACHED SHEET.)

Application Serial No.	Day/Month/Year Filed	abandoned
_____	_____	_____



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PATENT AND TRADEMARK OFFICE

I hereby appoint the following to prosecute and transact all business in the Patent and Trademark Office connected therewith:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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☒ SEE ATTACHED SHEET FOR SIMILAR INFORMATION AND SIGNATURE FOR ADDITIONAL JOINT INVENTORS.
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DECLARATION FOR USA PATENT APPLICATION

(including Design and PCT)

ADDITIONAL INFORMATION SHEET

(use as required)

Additional Prior Foreign Application(s):

Number	Country	Day/Month/Year Filed	Yes	No
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____

Additional 35 U.S.C. § 120 applications:

Application Serial No.	Day/Month/Year Filed	Status -- patented, pending, abandoned
_____	_____	_____
_____	_____	_____

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09666592-1451198

DECLARATION FOR USA PATENT APPLICATION

(including Design and PCT)

ADDITIONAL INFORMATION SHEET

(use as required)

Additional Prior Foreign Application(s):

Number	Country	Day/Month/Year Filed	Yes	No

Additional 35 U.S.C. § 120 applications:

Application Serial No.	Day/Month/Year Filed	Status -- patented, pending, abandoned

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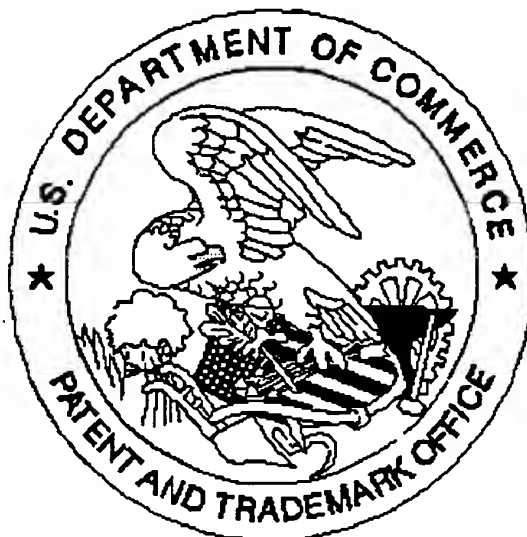
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